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(54) Amino acids in the treatment of neurodegenerative disorders.

(57) A composition for preventing in mammals the progressive exacerbation of symptoms associated with neurodegenerative disorders characterized by abnormal glutamate metabolism, elevated levels of peripheral glutamate or altered glutamate receptor characteristics includes amino acids that are effective in retarding the development of symptoms associated with neurodegenerative disorders, presumably by activating the enzyme glutamate dehydrogenase. In the preferred embodiment, the branched chain amino acids L-leucine, L-isoleucine, and L-valine are disclosed as effective therapeutic agents for the treatment of amyotrophic lateral sclerosis.

EP 0 341 895 A1

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AMINO ACIDS IN THE TREATMENT OF NEURODEGENERATIVE DISORDERS

This invention relates to compositions useful for treating neurodegenerative disorders in mammals. Specifically, compositions which are comprised of branched chain amino acids, such as L-leucine, L-isoleucine, and L-valine and which activate the enzyme glutamate dehydrogenase are disclosed. This invention also relates to protocols for their use, which are efficacious for retarding the progression of neurodegenerative disorders such as amyotrophic lateral sclerosis (ALS). The compositions disclosed have application for the treatment of neurodegenerative disorders characterized either by altered metabolism of glutamate, increased levels of peripheral glutamate, or altered glutamate receptor characteristics. The compositions disclosed presumably provide protection against the excitotoxic actions of glutamate.

The neurotoxicity of excitatory neurotransmitters has led to the development of the excitotoxicity hypothesis for neuronal cell death. [For review see, B. Engelsen, "Neurotransmitter Glutamate: its Clinical Importance", Acta Neurol Scand, 1986, 74:337-355; R. Schwarcz and B. Meldrum, "Excitatory Aminoacid Antagonists Provide a Therapeutic Approach to Neurologic Disorders", Lancet, 1985, 8447: 140-143] According to this hypothesis, sustained depolarization of neurons expressing receptors for excitatory amino acids ultimately leads to irreversible damage and their death. Sustained depolarizations may arise from several different mechanisms including increased release of glutamate, decreased reuptake of glutamate, or alterations in the characteristics of the glutamate receptor. The presence of one or more of these mechanisms in an individual may result in neurodegeneration and the onset of clinical symptoms. Several neurodegenerative disorders have been identified which are associated with one or more of the above alterations of glutamate activity. [B. Engelsen, "Neurotransmitter Glutamate: its Clinical Importance", Acta Neurol Scand, 1986, 74:337-355. The contents of this reference are herein incorporated by reference].

As toxicity has been associated with sustained depolarization through receptor activation, the predominant approach towards preventing the neurotoxic effect of neuroexcitatory neurotransmitters has been to develop effective antagonists for neuroexcitatory receptors. [R. Schwarcz and B. Meldrum, "Excitatory Aminoacid Antagonists Provide a Therapeutic Approach to Neurologic Disorders", Lancet, 1985, 8447: 140-143; S. M. Ross, M. Seelig, and P. Spencer, "Specific Antagonism of Excitotoxic action of 'Uncommon' amino acids in Organotypic Mouse Cortical cultures," Brain Res., 1987, 425:120-127] An example of this approach is the study of Guam ALS. The neurotoxic acitivity of beta-N-Methylamino-L-alanine (BMAA) and beta-N-oxalylamino-L-alanine (BOAA), com pounds which are present in seeds of Cycas circinalis and Lathyrus sativus, respectively, and which have been implicated in the etiology of Guam ALS, have been selectively antagonized in a concentration dependent manner by compounds which are specific antagonists of glutamate receptor sites. Thus, the development of specifically designed antagonists, especially those which block NMDA glutamate receptors, has been suggested to be "[o]ne of the most promising concepts concerning a rational treatment of neurodegenerative disorders[.]" [R. Schwarcz and B. Meldrum, "Excitatory Aminoacid Antagonists Provide a Therapeutic Approach to Neurologic Disorders, Lancet, 1985, 8447: 140-143]

Amyotrophic lateral sclerosis is a relentlessly progressive neurological disorder that is manifested primarily by muscle weakness, wasting, and spasticity, usually resulting in death from debilitating disease in 2-5 years [Brain, The Lord, Croft P., Wilkinson M. The course and outcome of motor neuron disease. In: Norris FH, Jurland LT, eds. Motor Neuron Diseases. New York: Grune and Stratton, 1967:22]. No treatment is known at present that can influence the course of the disease.

The cause of ALS is unknown. Recent studies [Plaitakis A., Caroscio JT. "Abnormal glutamate metabolism in amyotrophic lateral sclerosis, " Ann Neurol 1987. 22:575-579], however, have shown that the metabolism of the neuroexcitotoxic amino acid glutamate [Olney JW. "Neurotoxicity of excitatory amino acids", In: McGeer EG, Olney Jw, McGeer PL eds. Kainic Acid as a Tool in Neurobiology. New York: Raven Press, 1978: 95-121] is altered in this disorder and that the possibility exists that neurodegenerative processes in ALS may be mediated by neuroexcitotoxic mechanisms [Plaitakis A. Caroscio JT. "Abnormal glutamate metabolism in amyotrophic lateral sclerosis," Ann Neurol 1987. 22:575-579; Spencer PS, Nunn PB, Hugon J. et al. "Guam amyotrophic lateral sclerosis-Parkinsonism-dementia linked to a plant excitant neurotoxin," Science, 1987, 237:517-522].

Abnormal glutamate metabolism has, in addition to ALS, been shown to occur in patients with late-onset multi-system atrophic disorders associated with decreased activity of the enzyme glutamate dehydrogenase (GDH). [Plaitakis A., Berl S., Yahr MD. "Abnormal glutamate metabolism in an adult-onset neurological disorder," Science, 1982, 215:193-196]. Some of these patients with decreased GDH activity have been described to show predominant motor neuron involvement, or atypical ALS. [Plaitakis A., Berl S., Yahr MD. "Neurological disorders associated with deficiency of glutamate dehydrogenase", Ann. Neurol. 1984, 15 :

144-153] GDH is particularly important in the biology of the nervous system and the integrity of motor neurons. GDH can be activated by the branched chain amino acids (BCAA) L-leucine and L-isoleucine [Yielding KL, Tomkins GM. "An effect of L-leucine and other essential amino acids on the structure and activity of glutamate dehydrogenase," Proc. Natl. Acad. Sci. USA, 1961, 47:983].

5 This invention relates to the use of dietary supplementation with effective amino acid(s), especially with the branched chain amino acids (BCCAs) L-leucine, L-isoleucine and L-valine to benefit patients with idiopathic ALS. These compounds have been shown to allosterically activate, *in vitro*, glutamate dehydrogenase [Yielding KL, Tomkins GM. "An Effect of L-leucine and other essential amino acids on the structure and activity of glutamate dehydrogenase," Proc. Natl. Acad. Sci. USA, 1961, 47:983] and modify 10 the metabolism of glutamate [Dennis S., Clark, J. B. "The synthesis of glutamate by rat brain mitochondria," J. Neurochem 1986; 46:1811-1819], which is reported to be altered in ALS [Plaitakis A., Caroscio J. T. "Abnormal glutamate metabolism in amyotrophic lateral sclerosis," Ann Neurol 1987. 22:575-579]

A recent report describes the use of L-threonine to modify the progression of ALS. [B. M. Patten and L. M. Klein, "L-Threonine and the Modification of ALS," Neurology, 1988, 38 (Suppl 1) :354-355] To substitute 15 for L-threonine, L-leucine or L-isoleucine were used as placebos. Patients treated with placebo were reported to "revert to previous status."

This invention relates to compositions for treating neurodegenerative disorders, at least those which are not the result of a defect in the enzyme glutamate dehydrogenase.

The preferred compositions are comprised of three branched chain amino acids (BCAA) which may be 20 administered to patients in therapeutically effective amounts.

Twenty-two patients with amyotrophic lateral sclerosis (ALS) were entered into a double-blind, randomized, placebo-controlled trial to test whether the administration of branched-chain amino acids could favorably influence the course of the disease. Eleven patients received daily 12 grams of L-leucine, 8 grams of L-isoleucine and 6.4 grams of L-valine, given orally, and the remainder received placebo.

25 During the one-year trial, the placebo-treated patients showed a linear decline in their functional status consistent with the natural history of the disease. In contrast, the amino acid-treated patients showed a substantial stabilization of their neurologic status with significantly better main tenance of extremity muscle strength and continued ability to remain ambulatory than those treated with placebo.

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BRIEF DESCRIPTION OF DRAWINGS

Figure 1A. Mean Spinal Scores in the Branched-Chain Amino Acid (Treated) and Placebo Groups. (N) 35 is the number of patients who were clinically assessed at each time point as described in the Methods. The (N) decreased during the progress of the trial due to 1) death from the disease or severe disability precluding a visit to the clinic, 2) withdrawal from the study and 3) occasionally, non-compliance with the clinic appointments (Table 3). Two amino acid treated patients who could not attend the clinic at 12 months were instead evaluated at the 14th month of the trial. Values are means for each group \pm SEM.

40 Symbols refer to comparisons between the placebo and the amino acid treated group at each time point (using the unpaired t test). * Denotes p<0.02., † p<0.05 and ‡ p<0.1 of placebo compared to treated group.

Figure 1B. Comparison of spinal score differences from baseline (Δ spinal score) for placebo and amino acid-treated (treated) groups. The difference was calculated by subtracting the baseline value from the value at each time point for each subject. Values are means \pm SEM. For explanation of symbols set Fig. 45 1A.

Figure 2. Comparisons of the spinal score differences from baseline for only those amino acid-treated (treated) and Placebo-treated (placebo) patients who had complete observations for 12 months. Repeated measures analysis of variance showed that the time by group interaction was significant at p = 0.033 (F = 3.33) level.

50 Figure 3. Changes in Bulbar Scores for the Amino Acid (treated) and Placebo Groups. Bulbar function was clinically assessed at each time point as described in the methods. Values are means for each group \pm SEM and represent differences from baseline values (Δ bulbar score) as described in Fig. 1B.

Figure 4. Results of the branched-chain amino acid oral test dose. After overnight fasting, the ALS patients randomly receiving treatment received orally 3.0 grams of L-leucine, 2.0 grams of L-isoleucine and 55 1.6 grams of L-valine mixed thoroughly in dietetic jelly. Blood samples were drawn at the times indicated and processed as previously described(5). Data points represent mean values from 9 patients \pm SEM. * Denotes p<0.001 and † p<0.005. Symbols refer comparisons to 0 time values according to the student's t-test.

DETAILED DESCRIPTION OF THE INVENTION

5 In accordance with the subject of this invention a treatment is disclosed which is efficacious for preventing the progressive exacerbation of symptoms associated with neurodegenerative disorders characterized by abnormal glutamate metabolism, elevated levels of peripheral glutamate, or altered glutamate receptor characteristics. The treatment is comprised of administering to individuals in need of treatment a mixture of amino acids which activate the enzyme glutamate dehydrogenase.

10 Glutamate has been implicated in the pathogenesis of several human neurodegenerative disorders. Various human neurodegenerative disorders have been associated with one or more alterations involving glutamate including altered metabolism of glutamate, increased peripheral levels of glutamate, or altered glutamate receptor characteristics.

15 Glutamate, is an excitatory neurotransmitter which is also toxic to neurons. The neurotoxic activity of several excitatory neurotransmitters has led to the development of "excitotoxic models for human diseases." [R. Schwarzs and B. Meldrum. "Excitatory Aminoacid Antagonists Provide a Therapeutic Approach to Neurological Disorder," *Lancet*, July 20, 1985, 140-143. The contents of which are herein incorporated by reference] GDH catalyzes the conversion of glutamate to alpha-ketoglutarate. The disclosed invention provides treatments and compositions which presumably increase the endogenous activity of GDH thereby increasing the catabolism of glutamate as well as modifying the glutamatergic transmission mechanisms in the CNS to decrease its toxic activity. This is a completely novel approach to the treatment of neurodegenerative disorders. Most other approaches to the treatment of neurodegenerative disorders arising from excitotoxic mechanisms have focused on the use of receptor antagonists to inhibit the action of excitatory neurotransmitters such as glutamate.

20 Thus, this invention relates to the treatment of disorders that have been associated with glutamate toxicity. Such disorders include, but are not limited to amyotrophic lateral sclerosis, Alzheimer's disease, Huntington's disease, spinocerebellar degenerations, and other disorders characterized by degeneration of motor and other neurons of the central nervous system. This invention may have application as well and be useful for the treatment of other types of neurological disorders in which glutamate may be involved, such as seizure disorders, ischemic brain injury, or stroke, multiple sclerosis, and AIDS. This invention may have veterinary application as well.

25 The treatment comprises a composition comprised of one or more amino acids. Amino acids which activate glutamate dehydrogenase or are effective, either alone or in combination with other amino acids, for the treatment of one or more neurodegenerative disorders are appropriate ingredients of the composition. Such compositions may include branched chain amino acids. A preferred composition is comprised of L-leucine, L-isoleucine, and L-valine. The preferred composition has particular application to the treatment of amyotrophic lateral sclerosis (ALS).

30 Compositions provided by this invention should be administered so as to maintain minimum effective therapeutic concentrations of the composition constituents in plasma and other appropriate tissues. A preferred composition is comprised of L-leucine, L-isoleucine and L-valine, wherein the ratio of the amounts of L-leucine: L-isoleucine:L-valine present in the composition range from 0.3-30:0.2-20:0.16-16. The preferred range for the ratio of the amount of L-leucine:L-isoleucine: L-valine present in the composition is 3.0:2.0:1.6. For the treatment of neurodegenerative disorders, and particularly ALS, the amino acid composition should provide a total daily dose per individual of about 0.3 to 30 grams of L-leucine, 0.2 to 20 grams of L-isoleucine, and 0.16 to 16 grams L-valine. The preferred ranges of the total daily dose of each of the amino acids of the composition are about: 12 to 18 grams of L-leucine, 8 to 12 grams of L-isoleucine, and 3.2 to 9.6 grams of L-valine. The total daily dose may be divided so as to be administered using a variety of different protocols. Preferred protocols provide for the administration of the composition four or six times per day in amounts such that the total amount of composition administered per day is equal to the desired total daily dose. Administration of the composition may either be orally or parenterally.

35 Plasma levels of the amino acids of the composition may be monitored to assess whether effective levels are being maintained.

The doses of amino acids are sufficient to increase plasma levels above untreated levels. The ranges of the percent increase over untreated levels for each of the amino acids are about 50 to 600 % for L-leucine; 40 to 500 % for L-isoleucine; and 25 to 300 % for L-valine. The preferred ranges for the percent increase in plasma levels over untreated levels are about 100 to 300 for L-leucine, 80 to 250 % for L-isoleucine, and 50 to 150 % for L-valine.

EXAMPLE 1

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Patients and MethodsPatients

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Twenty-two patients were studied. The criteria for entry included (1) unequivocal diagnosis of ALS after a complete clinical investigation and EMG confirmation, (2) a stage of disease in which all patients were ambulatory at the time of entry, (3) normal biochemical and hematological profile and (4) lack of evidence for a systemic disease.

15

The patients were entered into the study sequentially as seen in the Mt. Sinai Hospital, New York, ALS clinic and randomly assigned to receive either BCAA powder or a similar amount of placebo powder (starch), with care taken to have equal numbers of subjects in both study groups. The BCAA powder consisted of 3.0 grams of L-leucine, 2.0 grams L-isoleucine and 1.6 grams of L-valine. Patients were instructed to take this orally four times daily between meals throughout the entire twelve month study period.

20

Each patient was then seen at three-month intervals for the clinical and laboratory assessments described below. The family, the patient, the nursing staff and the neurologists who performed the clinical evaluation were blinded as to the medication being used. The trial was terminated and the code was broken after all patients had completed twelve months on treatment.

25

Administration of L-leucine alone has been found to severely depress the plasma and CSF levels of the other two BCAs; however, when given in tandem with them, it produces significant increases in blood and CSF levels of all three compounds [Plaitakis A., Berl S., Yahr MD. "The treatment of GDH-deficient olivopontocerebellar atrophy with branched chain amino acids," Neurology 1983; 33 : (suppl 2) : 78 which is hereby incorporated by reference].

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Neurologic Evaluation

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Table 1.

Method of Estimating Bulbar and Spinal Scores

5 (a) Bulbar score:

Normal = 3
Mild impairment = 2
Moderate impairment = 1
Marked impairment = 0

10 Evaluation by this scale was made for
the following bulbar functions:

- 15 1) facial movement
2) tongue movement
3) palatal movement
4) dysarthria
20 5) dysphagia

Maximum score = 15 (All bulbar
functions were weighted equally)

25 (b) Spinal score

A power rating for individual muscle
groups was obtained by the following
scale;

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5 = normal strength
 4.5 = able to overcome marked resistance
 4 = able to overcome moderate resistance
 3.5 = able to overcome mild resistance
 3 = able to overcome gravity but not resistance
 10 2 = able to contract muscle with gravity eliminated
 1 = flicker of movement
 0 = no movement

15 Evaluation by this scale was made for the following paired (right and left) muscle groups:

- 1) arm abduction
- 2) elbow flexion
- 20 3) elbow extension
- 4) wrist flexion
- 5) wrist extension
- 6) finger flexion
- 7) finger extension
- 25 8) finger abduction
- 9) hip flexion
- 10) knee flexion
- 11) knee extension
- 12) foot dorsiflexion
- 30 13) foot plantar flexion

Maximum score = 130 (All muscle groups were weighted equally)

35

Laboratory Studies

40 Baseline blood chemistries and complete blood counts (CBC) were obtained. To assure bioavailability of the BCAA, an oral test dose of the above-described mixture or placebo was given after overnight fasting at baseline. Blood samples were drawn at 0, 60 and 120 minutes after dosing for multi-chemistries, CBSs and serum ammonia and amino acid levels. In addition, a random blood sample was obtained for the above studies at three month intervals with no attention paid to time of oral dosing. Plasma amino acid levels were assayed as previously described. [A. Platakis and J. Caroscio, "Abnormal Glutamate Metabolism in 45 Amyotrophic Lateral Sclerosis," Ann Neurol, 1987, 22:575-579. The contents of this article are herein incorporated by reference]

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Mean clinical scores for the two groups were calculated for values at baseline and at each three-month interval and were compared using the unpaired t-test. In addition the differences between the baseline and the scores obtained in each three month interval were statistically compared for the two groups using again the unpaired t-test. All t-tests were two sided.

55 Repeated measures analysis of variance was calculated on the clinical scores for the time periods 0, 3, 6, 9, and 12 months for all patients having complete data. The same analysis was also done for the time periods 0, 3, 6, and 9 months only, as well as for 0, 3, and 6 months only.

The outcome of the study was evaluated using two criteria: (1) rate of decline in scores of each group

and (2) continued ability to remain ambulatory at termination of the study. The proportion of success of the second criterion in each of the two groups was statistically compared using the chi-squared statistic and Fisher's exact test. Included in this analysis were all patients who completed the trial as well as those who, while under treatment, progressed to the point of wheelchair confinement (classified as failures) but did not complete one year on the trial either due to death or subsequent withdrawal. Patients who withdrew while still ambulatory were excluded from analysis because they could not be classified as successes without having completed 12 months of observation.

10

Results

Twenty-two patients meeting the requirements for entry were included in the study. Eleven patients were assigned to BCAA treatment, the remainder to placebo. An additional patient assigned to BCAA was lost to follow-up within a month and was not included in any of the analyses. The clinical characteristics of the two treatment groups did not differ significantly (Table 2).

20

Table 2.

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Clinical Characteristics of ALS Patients at Entry into the Study, According to Treatment Group			
Characteristics		Branched-chain Amino-Acids	Placebo
Male		10	9
	Female	1	2
Disease Duration			
< 2 years		7	6
	> 2 years	4	5
Mean Disease Duration (years)		1.8±2.0*	1.9±1.2*
Mean Age (years)		48.5±11.9*	53.7±11.5*
Mean Spinal Score (Normal = 130)		115.9±15.6*	114.7±11.1*
Mean Bulbar Score (Normal = 15)		12.5±2.2*	13.2±2.2*

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* Values are means±S.D.

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Table 3 shows the number of patients who remained on the protocol throughout the trial, as well as those who withdrew, became respirator-dependent or died of their disease.

Table 3

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Number of ALS Patients On And Off Protocol During The Trial				
Time on the Trial	Amino Acid Treated		Placebo Treated	
	On Protocol	Off Protocol	On Protocol	Off Protocol
0	11	0	11	0
3 months	11*	0	11	0
6 months	10*	1 (withdrawal)	10*	1 (withdrawal)
9 months	8*	2 (withdrawals)	9	1 (death)
12±2 months	8	0	6	3 (death: 1 withdrawal: 1 respirator: 1

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* One patient was unable to attend clinic for evaluation at this time but remained on treatment and was evaluated on subsequent visits.

Fig. 1A shows the mean spinal scores obtained in all patients in each group at three-month intervals and Fig. 1B shows the changes in these scores (values at the time points minus baseline values). Fig. 2 shows changes in spinal scores of only those patients who had complete observations for 12 months. The changes were significantly worse in the placebo-treated patients than in the BCAA-treated patients.

Repeated measures analysis of the spinal scores for all patients who completed the trial revealed significant differences in the rate of progression of the disease between the two groups. Thus, a statistically significant time X group interaction was found for the time periods: (a) 0-6 months ($F = 7.13$, $p = 0.0003$) ($N = 17$, drug group = 9, placebo group = 8) and (b) 0-9 months ($F = 5.17$, $p = 0.0004$, $N = 15$, drug group = 7, placebo group = 8) and (c) 0-12 months ($F = 3.6$, $p = 0.013$) ($N = 12$, drug group = 7, placebo group = 5).

Due to progressive decline, five of the nine placebo-treated patients lost their ambulation during the trial with two subsequently dying and another becoming respirator-dependent (Table 3). In contrast, only one of the nine BCAA-treated subjects lost ambulation during observation ($p < 0.05$ by the chi square; $p = 0.066$ by the Fisher's exact test). Two patients in each group who withdrew (Table 3) while still ambulatory were not included in this analysis (see Methods).

In contrast to the spinal scores, the changes in the bulbar scores were not significantly different between the two treated groups (Fig. 3). Also, there was no significant time X group interaction with respect to the bulbar scores. However, there was a significant change in the bulbar scores overall for baseline vs. nine months ($p = 0.025$, $F = 6.79$) and baseline vs. 12 months ($p = 0.016$, $F = 8.16$).

Plasma levels of leucine, isoleucine and valine showed marked increases following oral dosing (Fig. 4). Levels of leucine increased by 300% and 292%, of isoleucine by 393% and 283% and of valine by 95% and 53% after 1 and 2 hours, respectively. Routine blood chemistries and complete blood counts did not show any abnormalities following these oral loadings, except for mild transient elevations in the plasma ammonia levels (data not shown). Analysis of blood samples obtained randomly at various intervals during the one-year trial revealed that the amino acid-treated patients maintained plasma levels of BCAA that were significantly higher than those of the placebo-treated patients (leucine by 112%, isoleucine by 93% and valine by 61%). Also, complete blood counts and blood chemistries (including ammonia levels) did not show significant changes in the patients who were treated chronically with the BCAA.

Long-term controlled trials have proven difficult to perform in ALS [Shafter SR, Olarte MR. "Methodological considerations for clinical trials in motor neuron disease." In: Rowland P ed. Human Motor Neuron Diseases, New York, Raven Press, 1982, pp. 559-568] because of the high rate of patient loss (due to death or discontinuation of treatment) which complicates the analysis and interpretation of the data. An attrition rate of study patients as high as 70% has been described in one-year, double-blind, controlled trials [Olarte MR, Shafter SR. "Levamisole is ineffective in the treatment of amyotrophic lateral sclerosis," Neurology 35:1063-1066].

Although some cases were lost during this study, the majority of treated patients completed the one-year trial. Analysis of the spinal scores, using all available measurements of every patient who was included in the trial, as well as the measurements of only those patients who had complete observations, revealed significant differences in the rate of progression between the two groups, regardless of the inclusion or exclusion of the patients who did not have complete data or did not complete the one-year trial.

Thus, the deterioration in the mean spinal performance scores for the BCAA-treated patients was 4% and 9% of the initial scores for 6 and 12 months respectively. In contrast, in the placebo group, deterioration of these scores was 18% and 38% for 6 and 12 months respectively. The decline in motor performance of our placebo-group was comparable to that described by Olarte et al. [Olarte MR, Shafter SR. "Levamisole is ineffective in the treatment of amyotrophic lateral sclerosis," Neurology, 1985, 35:1063-1066], who found a 15-25% and 20-30% decline in mean score for 6 and 12 months respectively, although direct comparisons are confounded by different scoring methods. Also, the death rate in the placebo group (2 out of 11 or 18.2%) was comparable to that in the study by Olarte et al. [Olarte MR, Shafter SR. "Levamisole is ineffective in the treatment of amyotrophic lateral sclerosis," Neurology, 1985, 35:1063-1066] (18.5% and 20.5% for their two groups).

The favorable results of this invention are clearly not due to difference in the severity of illness and approximate duration of illness prior to treatment in the two groups studied, which were rather well-matched. Because of the small sample size and because there is a substantial variation in the natural history of the disorder modifications of the treatment produced are likely to be developed based on the principle disclosed in this invention. Of some concern is the fact that no significant differences in the bulbar scores of the two groups were found. It should, however, be noted that clinical rating of bulbar motor function is less sensitive than spinal rating and that bulbar dysfunction tends to progress slower than spinal weakness in ALS patients.

This preliminary trial establishes a basis for a new therapeutic approach to ALS, and other disorders

characterized by altered glutamate metabolism.

Example 2

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The ability of L-leucine to activate, *in vitro*, GDH present in post mortem brain tissue obtained from control patients and patients with multisystem atrophy-multineuron disease was determined. (Table 4) Tissue preparation and GDH determinations were done according to methods previously reported. [A. D. Colon, A. Plaitakis, S. Berl, and D. D. Clark. "The purification and characterization of soluble and particulate Glutamate Dehydrogenase in Rat Brain," *Neurochemistry*, 1986, 46: 1811-1819. The contents of this publication are herein incorporated by reference] L-leucine produces a significant activation of GDH in tissue obtained from control cerebellum, and from tissue obtained from multisystem atrophy-multineuron degeneration disease cerebellum and frontal cortex. L-leucine concentrations as low as 0.15 mM are effective in producing increases in GDH activity and this effect is concentration dependent. The demonstration of leucine mediated activation of GDH at concentrations between 0.15 and 0.5 mM is important as plasma leucine levels of between about 0.45 to 0.30 mM at 1 and 2 hours, respectively, are obtained following administration of the amino acid composition to patients with ALS. Figure 4.

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Table 4

In Vitro Leucine Activation of Membrane Bound GDH in Purified Brain			
Leucine Concentration (mM)	Percent Baseline GDH Activity		
	Multisystem Atrophy Multineuron Disease		
	Control Patient	Patient 1	Patient 2
	Cerebellum	Cerebellum	Frontal Cortex
0.0	100	100	100
0.15	-	-	122
0.25	117	117	128
0.50	135	153	141
1.0	143	186	-
2.5	155	-	-
5.0	155	176	-
10	160	176	-

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The aforementioned articles are incorporated herein by reference.

While we have hereinbefore described a number of embodiments of this invention, it is apparent that the basic constructions can be altered to provide other embodiments which utilizes the compositions of this invention.

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Claims

1. A pharmaceutical composition to be used for the treatment of a neurodegenerative disorder which includes one or more amino acids which alone or in combination with each other are effective for treating one or more of said neurodegenerative disorders.

2. A pharmaceutical composition to be used for the treatment of a neurodegenerative disorder which includes one or more amino acids which alone or in combination with each other activate the enzyme glutamate dehydrogenase.

3. A pharmaceutical composition according to claim 1 or claim 2, wherein there is a mixture of amino acids which comprises branched chain amino acids.
4. A pharmaceutical composition according to claim 3, wherein the mixture of amino acids comprises all of L-leucine, L-isoleucine, and L-valine.
5. A pharmaceutical composition according to claim 4, wherein the ratio of the amount of L-leucine:L-isoleucine:L-valine present in the composition is in the ranges 0.3-30:-0.2-20:0.16-16.
6. A pharmaceutical composition according to claim 5, wherein the ratio of the amount of L-leucine:L-isoleucine:L-valine present in the composition is about 3.0:2.0:1.6.
7. A pharmaceutical composition according to any one of the preceding claims, wherein the neurodegenerative disorder to be treated is amyotrophic lateral sclerosis.
8. A pharmaceutical composition according to any one of claims 1 to 6 wherein the neurodegenerative disorder to be treated is alzheimer's disease, Huntington's chorea, spinocerebellar degenerations, or other disorder characterized by degeneration of motor and other neurons of the central nervous system.
9. A pharmaceutical composition according to any one of claims 4-6 in unit dosage form, the unit dosage being such as to give a patient a total daily dose within the range.
 - a. L-leucine, 0.3 to 30 grams;
 - b. L-isoleucine, 0.2 to 20 grams;
 - c. L-valine, 0.16 to 16 grams.
10. A pharmaceutical composition according to claim 9 wherein the total daily dose is within the range.
 - a. L-leucine, 12 to 18 grams;
 - b. L-isoleucine, 8 to 12 grams;
 - c. L-valine, 3.2 to 9.6 grams.
11. A pharmaceutical composition according to claim 9 or claim 10 wherein the unit is one quarter or one sixth of the total daily dose.
12. Use of a branched-chain amino acid in the manufacture of a pharmaceutical composition for the treatment of neurodegenerative disorders.
13. Use of a mixture of L-leucine, L-isoleucine and L-valine in the manufacture of pharmaceutical composition for the treatment of neurodegenerative disorders.

Claims for the following Contracting State: ES

1. A method of manufacturing a pharmaceutical composition to be used for the treatment of a neurodegenerative disorder which includes using in said manufacture one or more amino acids which alone or in combination with each other are effective for treating one or more of said neurodegenerative disorders.
2. A method of manufacturing a pharmaceutical composition to be used for the treatment of a neurodegenerative disorder which includes using in said manufacture one or more amino acids which alone or in combination with each other activate the enzyme glutamate dehydrogenase.
3. A method according to claim 1 or claim 2, wherein there is a mixture of amino acids which comprises branched chain amino acids.
4. A method according to claim 3, wherein the mixture of amino acids comprises all of L-leucine, L-isoleucine, and L-valine.
5. A method according to claim 4, wherein the ratio of the amount of L-leucine:L-isoleucine:L-valine present in the composition is in the ranges 0.3-30:-0.2-20:0.16-16.
6. A method according to claim 5, wherein the ratio of the amount of L-leucine:L-isoleucine:L-valine present in the composition is about 3.0:2.0:1.6.
7. A method according to any one of the preceding claims, wherein the neurodegenerative disorder to be treated is amyotrophic lateral sclerosis.
8. A method according to any one of claims 1 to 6 wherein the neurodegenerative disorder to be treated is alzheimer's disease, Huntington's chorea, spinocerebellar degenerations, or other disorder characterized by degeneration of motor and other neurons of the central nervous system.
9. A method according to any one of claims 4 to 6 in unit dosage form, the unit dosage being such as to give a patient a total daily dose within the range
 - a. L-leucine, 0.3 to 30 grams;
 - b. L-isoleucine, 0.2 to 20 grams;

c. L-valine, 0.16 to 16 grams.

10. A method according to claim 9 wherein the total daily dose is within the range

- a. L-leucine, 12 to 18 grams;
- 5 b. L-isoleucine, 8 to 12 grams;
- c. L-valine, 3.2 to 9.6 grams.

11. A method according to claim 9 or claim 10 wherein the unit is one quarter or one sixth of the total daily dose.

10 12. Use of a branched-chain amino acid in the manufacture of a pharmaceutical composition for the treatment of neurodegenerative disorders.

13. Use of a mixture of L-leucine, L-isoleucine and L-valine in the manufacture of a pharmaceutical composition for the treatment of neurodegenerative disorders.

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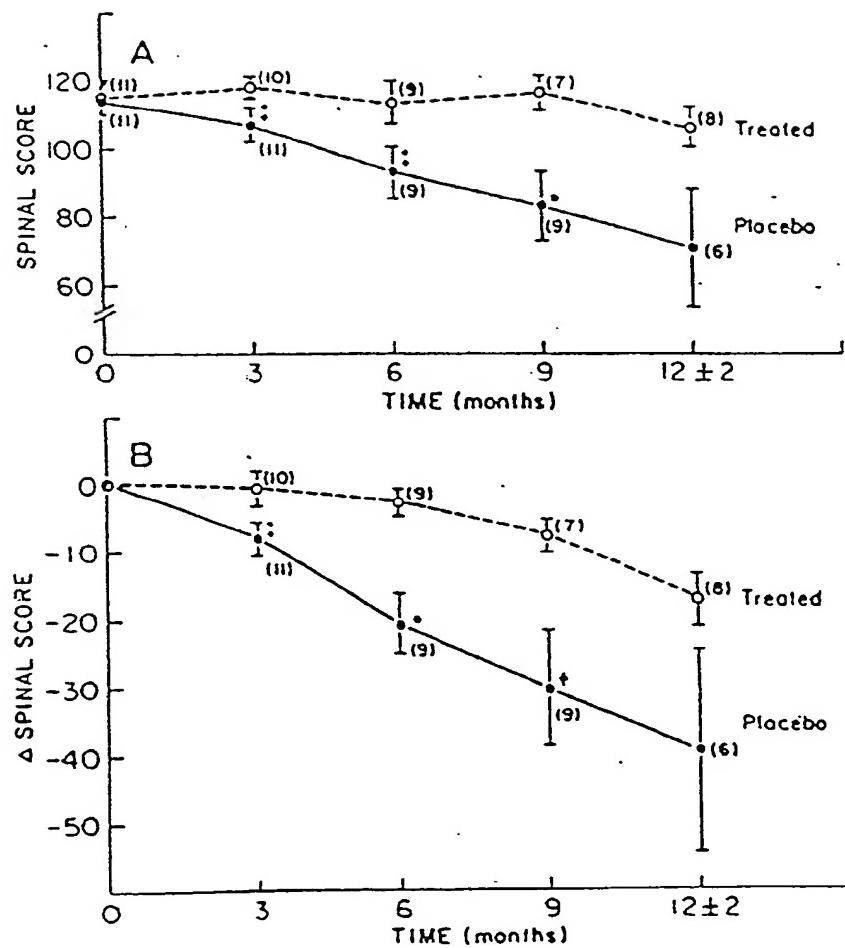


Fig. 1

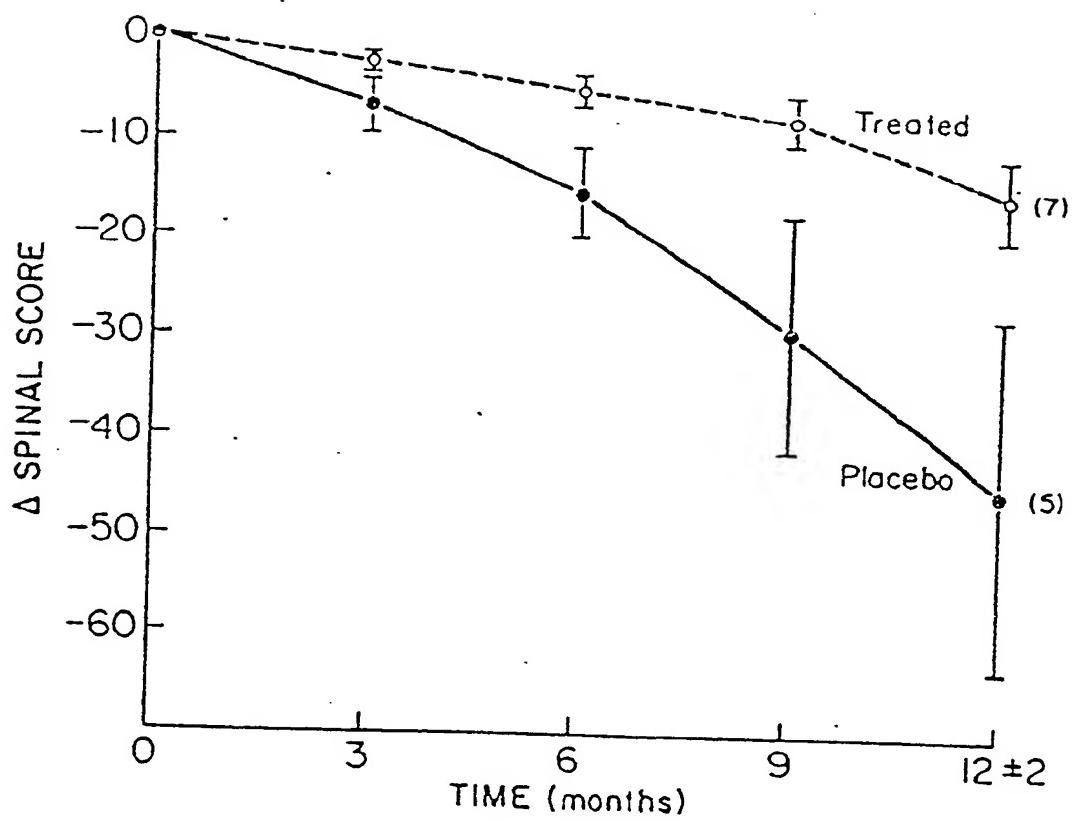


Fig. 2

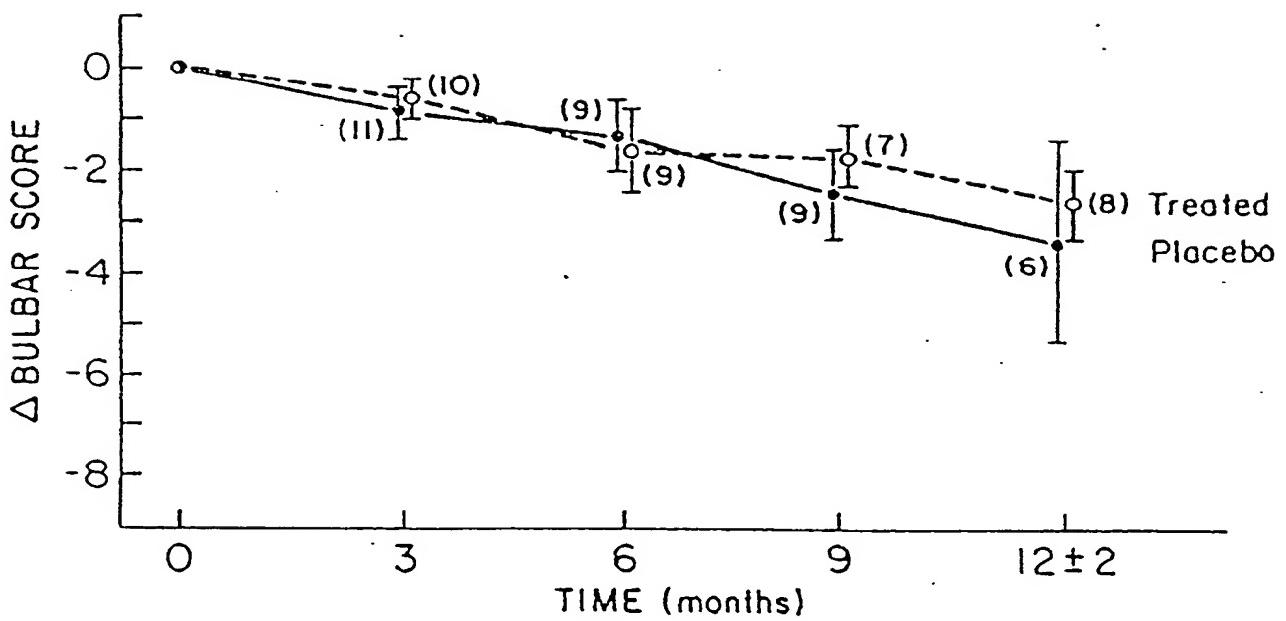


Fig. 3

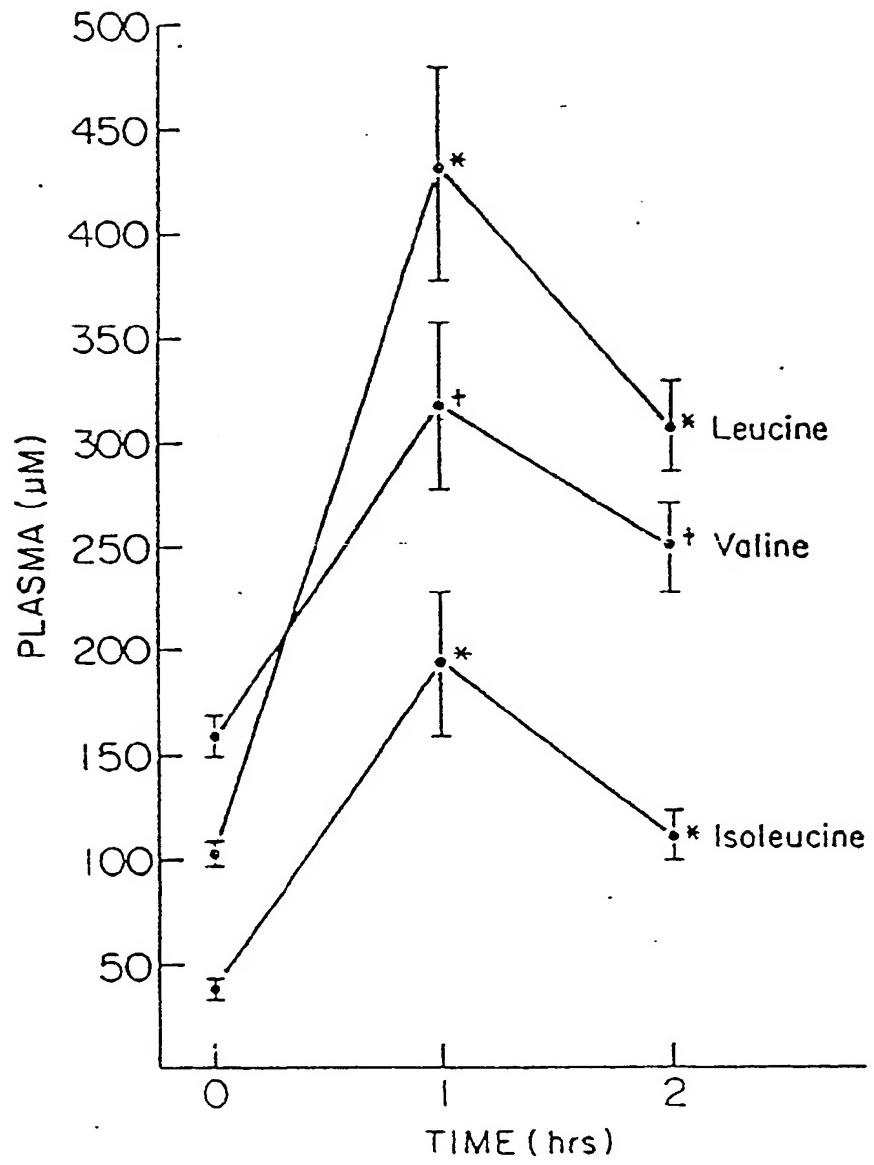


Fig. 4



EP 89 30 4463

DOCUMENTS CONSIDERED TO BE RELEVANT					
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)		
X	ROTE LISTE, no. 56 086, Editio Cantor, Aulendorf/Württ; "Lactostrict spezial" ---	1-13	A 61 K 31/195		
X	UNLISTED DRUGS, vol. 37, no. 5, May 1985, Chatham, New Jersey, US; "Ilevasorb" -----	1-13			
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int. Cl.4)		
			A 61 K		
Place of search	Date of completion of the search	Examiner			
THE HAGUE	25-08-1989	BRINKMANN C.			
CATEGORY OF CITED DOCUMENTS					
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document					
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document					

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